



# SmartPA Criteria Proposal

Drug/Drug Class:	Antiemetics, 5-HT3, NK1 and Other Select Non-Injectables PDL Edit	
First Implementation Date:	October 6, 2004	
Proposed Date:	December 16, 2021	
Prepared For:	MO HealthNet	
Prepared By:	MO HealthNet/Conduent	
Criteria Status:	□Existing Criteria	
	⊠Revision of Existing Criteria	
	□New Criteria	

# **Executive Summary**

Purpose: The MO HealthNet Pharmacy Program will implement a state-specific preferred drug list.

Why Issue Selected:

The 5-HT3 receptor antagonists are indicated for the prevention and treatment of nausea and vomiting associated with chemotherapy and radiotherapy. Diclegis® (doxylamine/pyridoxine) is indicated for the treatment of nausea and vomiting of pregnancy in women who do not respond to conservative management. Metoclopramide is indicated for diabetic gastroparesis and gastroesophageal reflux, as well as off-label use for treating nausea and vomiting associated with chemotherapy and radiation. Neurokinin-1 (NK1) receptor antagonists have indications that include prevention and treatment of nausea and vomiting associated with chemotherapy and radiotherapy and post-operative nausea and vomiting. The management of chemotherapy-induced nausea and vomiting is a critical aspect of treating cancer patients. The advent of agents within this therapeutic class was a significant breakthrough for the practice of oncology. However, because of the increased cost of these products, it is essential that therapy is appropriately monitored, and prudently utilized for the appropriate patient population.

Total program savings for the PDL classes will be regularly reviewed.

### Program-Specific Information:

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С	Preferred Agents	Non-Preferred Agents	
1:	Aprepitant Caps	Akynzeo <sup>®</sup> Caps	
	<ul> <li>Metoclopramide Soln/Tabs</li> </ul>	Anzemet®	
	<ul> <li>Ondansetron ODT/Soln/Tabs</li> </ul>	Bonjesta®	
		Diclegis®	
		Doxylamine/Pyridoxine	
		Emend® Caps/Pwd Packet	
		Gimoti®	
		Granisetron Tabs	
		Metoclopramide ODT	
		Reglan <sup>®</sup>	
		Sancuso®	
		Zofran <sup>®</sup> Tabs	
		Zuplenz <sup>®</sup>	

## SmartPA PDL Proposal Form

Type of Criteria:	<ul><li>☐ Increased risk of ADE</li><li>☐ Appropriate Indications</li></ul>	<ul><li>☑ Preferred Drug List</li><li>☐ Clinical Edit</li></ul>
Data Sources:	☐ Only Administrative Databases	□ Databases + Prescriber-Supplied

# **Setting & Population**

- Drug class for review: Antiemetics, 5-HT3, NK1 and Other Select Non-Injectables
- Age range: All appropriate MO HealthNet participants

# Approval Criteria

- For aprepitant capsules: maximum quantity of 3 doses per chemotherapy course OR
- Failure to achieve desired therapeutic outcomes with a trial on 1 or more preferred agents:
  - Documented trial period for preferred agents OR
  - Documented ADE/ADR to preferred agents AND
- For Diclegis and Bonjesta: adequate therapeutic trial of doxylamine (trial defined as 60/90 days)
- For Zuplenz: Clinical Consultant Review required for medical necessity of therapy

### **Denial Criteria**

- Therapy will be denied if all approval criteria are not met
- Lack of adequate trial on required preferred agents
- Claim exceeds maximum dosing limitations for the following:

Drug Description	Generic Equivalent	Max Dosing Limitations
ANZEMET 100 MG	DOLASETRON	1 tablet per day
ANZEMET 50 MG	DOLASETRON	1 tablet per day
GRANISETRON 1 MG	GRANISETRON	2 tablets per day
ZOFRAN 24 MG	ONDANSETRON	1 tablet per day
ZUPLENZ 4 MG	ONDANSETRON	3 films per day
ZUPLENZ 8 MG	ONDANSETRON	3 films per day

# ZUPLENZ 8 MG ONDANSETRON 3 films per day ZUPLENZ 8 MG ONDANSETRON 3 films per day Required Documentation Laboratory Results: Progress Notes: X MedWatch Form: Other: Disposition of Edit Denial: Exception Code "0160" (Preferred Drug List) Rule Type: PDL Default Approval Period

### 6 months

# References

- Evidence-Based Medicine and Fiscal Analysis: "Antiemetic Agents Therapeutic Class Review", Conduent Business Services, L.L.C., Richmond, VA; November 2021.
- Evidence-Based Medicine Analysis: "Antiemetics 5-HT3, THC Derivatives, NK1 & Other Agents", UMKC-DIC; September 2021.
- USPDI, Micromedex; 2021.
- Facts and Comparisons eAnswers (online); 2021 Clinical Drug Information, LLC.

